



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<p>(21) International Application Number: PCT/US91/05399 (22) International Filing Date: 31 July 1991 (31.07.91) (30) Priority data: 563,024 3 August 1990 (03.08.90) US (71) Applicant: EASTMAN KODAK COMPANY [US/US]; 343 State Street, Rochester, NY 14650 (US). (72) Inventor: YAU, Cheuk, Chung ; 313 Highridge Drive, Kingsport, TN 37660 (US). (74) Agent: STEVENS, John, F.; 343 State Street, Rochester, NY 14650-2201 (US).</p>		<p>(81) Designated States: AT (European patent), BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, KR, LU (European patent), NL (European patent), SE (European patent). Published With international search report.</p>
<p>(54) Title: POLYMER PELLET CONFIGURATION FOR SOLID-STATE POLYMERIZATION</p> <p>(57) Abstract</p> <p>Disclosed is a configuration for a polymer pellet having improved reactivity for solid-state polymerization and less tendency to stick during processing with heat. The pellets have a reduced path length for volatiles to travel to the exterior of the pellet, and reduced contact area in a packed condition.</p> <p style="text-align: center;">BEST AVAILABLE COPY</p>		

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⁺ It is not yet known for which States of the former Soviet Union any designation of the Soviet Union has effect.

POLYMER PELLET CONFIGURATION FOR
SOLID-STATE POLYMERIZATION

Technical Field

5 The present invention relates to polymer pellet
configuration for solid-state polymerization. More
specifically, the present invention relates to pellets
having a short path length for volatiles to be removed
from the pellets, and a reduced contact area among
10 pellets to minimize sticking of the pellets undergoing
solid-state polymerization in a stationary or moving bed
reactor.

 It is known that polyethylene terephthalate may be
prepared in the molten state by polycondensation
15 accompanied by liberation of ethylene glycol from
prepolycondensates in autoclaves of conventional design.
Intrinsic viscosities of approximately 0.7 are usually
obtained. Intrinsic viscosities as high as 1.00 can be
obtained in specifically designed reactors which improve
20 the evaporation of glycol. These prior art processes
have the disadvantage that high reaction temperatures,
high catalyst concentrations and long reaction time
periods are required to achieve high relative
viscosities. The inevitable result is an increasing
25 thermal degradation of the melt which manifests itself
by a large number of carboxyl end groups and an
intensive yellow coloring of the end product. A high
percentage of carboxyl end groups leads to strong signs
of degradation during the remelting of the
30 polycondensate. The discolorations are particularly
undesirable if pellucid polycondensates are needed for
the manufacture of, for instance, packing material for
foodstuffs or beverages. In addition, simultaneous with
the discoloration, elevated concentrations of aldehydes
35 are to be expected in the melt. These properties are

particularly undesirable if the end product is used to manufacture beverage bottles. The aldehydes diffuse into the liquid and affect the flavor of these beverages. Further, such aldehyde containing beverages
5 may be considered to be unfit for human consumption by certain regulatory bodies.

In order to avoid these disadvantages, it is known to produce high viscosity polyesters in the solid phase at temperatures below the melting point of these
10 condensates and under an inert gas blanket or vacuum. The drawbacks of this improvement are the large efforts and expenditure required to prevent the granulate from sticking in the course of the solid state polycondensation.

15 In one attempted solution to solve this sticking problem, the prior art has subjected the granulate to a continuous mechanically produced movement during the solid state polycondensation. Another prior art attempt to solve this stickiness problem is to effect the solid
20 state polycondensation in a fluidized bed in the presence of hot inert gases, such as nitrogen. In order to avoid agglomerations due to stickiness the prior art has also added the so-called anticaking agents to the surface of the polycondensed granulate.

25

Brief Description of the Drawings

Figure 1 is a plan view of a pellet according to the present invention taken along lines 1-1 of Figure 2; Figure 2 is a side view of the pellet shown in Figure 1;
30 Figure 3 is a modification of the pellet according to the present invention; and Figure 4 is a view showing two pellets stacked together.

Description of the Invention

According to the present invention, there is provided a polymer pellet having improved reactivity for solid state polymerization and less tendency to stick during processing with heat, the pellet having opposite faces generally lying in parallel planes, and in cross-section in a plane parallel to the faces the pellet having a generally rectangular central section, the corners thereof protruding into generally rounded lobes, the generally rectangular central section having dimensions A and B wherein the ratio of A to B is 1.2 to 6 and the ratio of B to the average radius of said lobes is 0.3 to 5.

The polymer of the pellet according to this invention may be any polymer which is to be polymerized in the solid state. Preferred polymers are polyesters, and in particular poly(ethylene terephthalate) or copolymers thereof.

Typically, polyesters such as poly(ethylene terephthalate) are produced to an intermediate molecular weight of, say 0.4-0.65 in a melt phase, by reacting a dicarboxylic acid or ester with a glycol using suitable catalysts under conventional reaction condition well known in the art. Following melt phase polymerization, the polymer may be subjected to further polymerization in the solid state.

Solid state polymerization is a well known technique for building up molecular weight of a polymer in the solid state. One conventional solid-stating operation takes place in so-called plug flow reactors wherein polymer pellets are introduced into the top of a tall cylindrical vessel and removed from the bottom at the same rate. During the residence times in the reactor, which is commonly in the order of 8-18 hours, the pellets reach high temperatures, generally

200-250°C. Molecular weight, as measured by inherent viscosity (I.V.), increases during the time the pellets are in the reactor. Typically, I.V. is increased from a range of 0.4-0.65 to greater than 0.7. This process is
5 thoroughly described in U.S. Patent No. 4,064,112, which is incorporated herein by reference.

Described herein is a design of the cross section of the pellets that incorporates into a pellet of a given weight a reduced path length the volatiles need to
10 travel to the exterior of the pellet to increase solid-state polymerization rate, and reduced contact area among pellets to minimize sticking. This design will also reduce the initial degradation experienced by the pellets during solid-state polymerization because
15 the shorter path allows the moisture content of the pellets to leave in much shorter times and minimize the probability of hydrolytic degradation. The design is shown in Figures 1 and 2.

Figure 1 illustrates pellet 10 having a generally rectangular central section 12 having dimensions A and
20 B. The corners of section 12, 14, 16, 18 and 20 are illustrated in broken lines and protrude into generally rounded lobes 22, 24, 26 and 28, respectively. The lobes are preferably rounded as shown in Figure 1, but
25 may be of the general shape shown in Figure 3 wherein the C dimension would be an average distance or radius as indicated in Figure 3 by broken line 30.

The optimal design has the ratio of B to C of 1 and the ratio of A to B ranging from 1.6 to 4, although the
30 benefit of this design can still be realized when the ratio of B to C varies from 0.3 to 5 and the ratio A to B varies from 1.2 to 6. When the ratio of B to C is greater than 1, the cross section of the pellet has the general shape as shown in Figure 1.

The advantage of the mentioned design can be demonstrated in Figure 4 wherein the pellets 10 and 11 can be stacked with minimum amount of surface contact. The added space between the two pellets also allows carrier gas to pass through to remove volatiles generated during solid-state polymerization thus increasing the rate of solid-state polymerization.

The pellets of the present invention can be obtained by conventional methods, such as by extruding the polymer melt through a die with the die holes taking the shape as shown in Figure 1. The extruded polymer can then be chilled to form a solid rod and be cut into pellets of desired lengths. The length (1) of the pellets preferably ranges from 0.05 to 0.2 inch but can vary from 0.03 to 0.3 inch. The dies can have any number of holes normally ranging from 1 to 100. The opposite upper and lower faces 13 and 15 of pellet 10 are theoretically in generally parallel planes. However, under production conditions, these faces may not be perfectly flat or parallel due to the fact that the cutter is cutting a continuously moving strand of polymer and upon cooling of the pellet there may be slight distortion.

The preferred design is for lobes 22, 24, 26 and 28 to be generally rounded. However, they do not have to be perfectly round and may even resemble portions of a square or rectangle as shown in Figure 3. In this case, the average diameter, or an arc 30 would be used as the C dimension.

The following example is submitted for a better understanding of this invention.

Pellets with cross sections taking the shape of a circle, square, rectangle and design according to this invention are prepared by extruding poly(ethylene) terephthalate with a 0.76 I.V. through a die plate with

the corresponding shape of die holes. The die holes have the following dimensions: circular, 5.09 mm in diameter; square, 4.51 x 4.51 mm; rectangular, 2.61 x 7.82 mm; according to invention, A = 6.39, B = 2.13, C = 1.69 mm (dimensions from Figure 1). The circular, square and rectangular shapes are for comparison. The extruded rods are quenched in chilled water and cut into pellets with an average weight of 0.0177 g/pellet using a rotary cutter. The pellets are crystallized at 180°C and the solid-state polymerization rates are measured. The results are shown in Table 1.

Table 1. Effect of Pellet Shape on Solid-State Polymerization Rate of PET

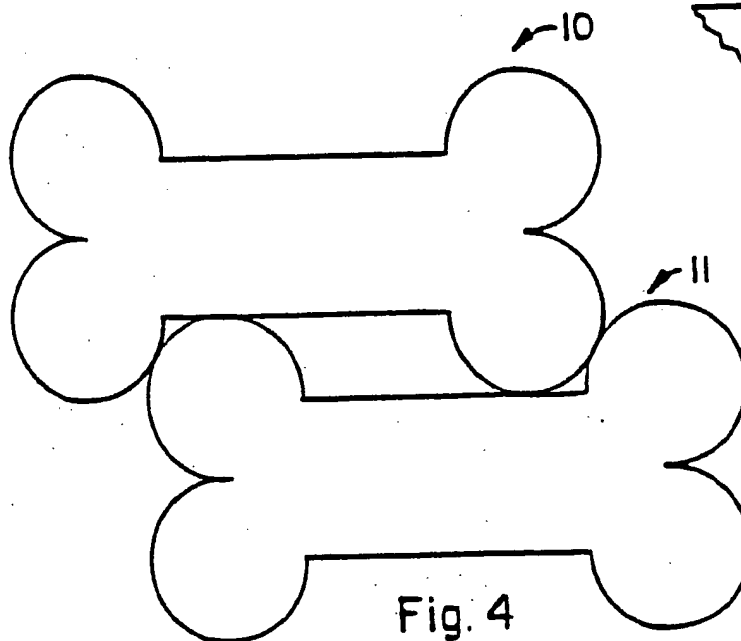
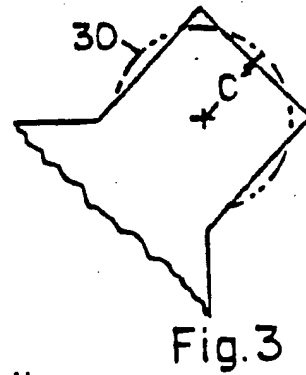
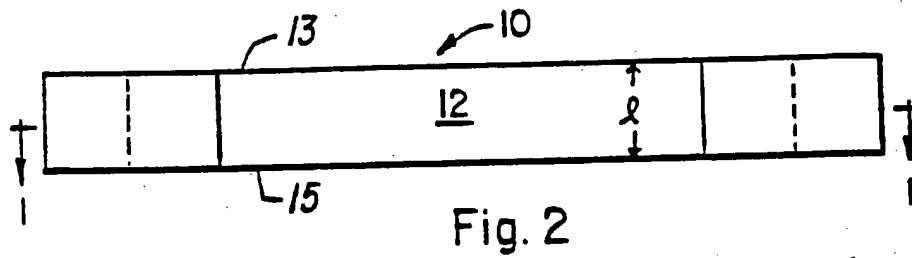
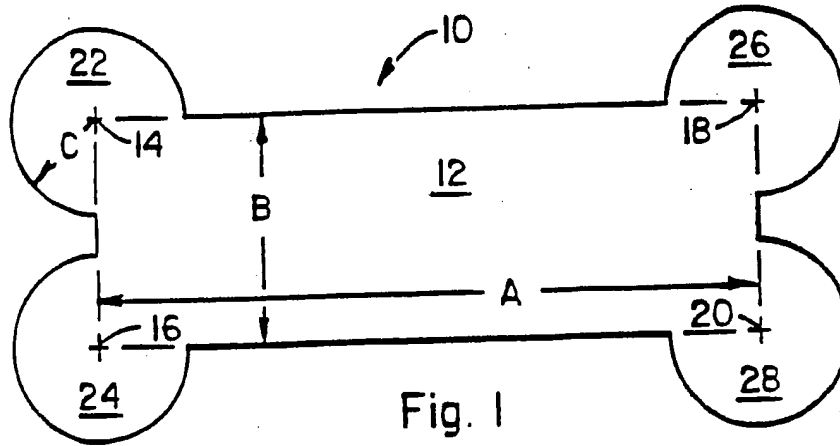
15	<u>Pellet Shape</u>	<u>Solid-State Polymerization Rate (dL/g-h)</u>
20	round (comparative)	0.01410
	square (comparative)	0.01372
	rectangle (comparative)	0.01594
	Design shown in Figures 1 and 2	0.01934

25 I.V. (inherent viscosity) is measured at 25°C using 0.50 gram of polymer per 100 mL of a solvent consisting of 60% by weight phenol and 40% by weight tetrachloroethane.

30 The invention has been described in detail with particular reference to preferred embodiments thereof, but it will be understood that variations and modifications can be effected within the spirit and scope of the invention.

CLAIMS

1. A polymer pellet having improved reactivity for
solid-state polymerization, and less tendency to
stick during processing with heat, said pellet
characterized by having opposite faces lying
generally in parallel planes, and in cross-section
in a plane parallel to said faces said pellet
having a generally rectangular central section, the
corners thereof protruding into lobes, said
generally rectangular central section having
dimensions A and B wherein the ratio of A to B is
1.2 to 6 and the ratio of B to the average radius
of said lobes is 0.3 to 5.
2. A polymer pellet according to Claim 1 wherein said
corners protrude into generally rounded lobes.
3. A polymer pellet according to Claim 1 wherein the
ratio of A to B is 1.6 to 4.
4. A polymer pellet according to Claim 1 wherein the
ratio of B to C is 0.7 to 3.
5. A polymer pellet according to Claim 1 wherein the
length of the pellets is 0.05 to 0.2 inch.
6. A polymer pellet according to Claim 1 wherein the
polymer is a polyester.



INTERNATIONAL SEARCH REPORT

International Application No PCT/US 91/05399

I. CLASSIFICATION OF SUBJECT MATTER (If several classification symbols apply, indicate all) ⁶ According to International Patent Classification (IPC) or to both National Classification and IPC IPC5: C 08 G 63/80, C 08 J 3/12														
II. FIELDS SEARCHED <div style="text-align: center; border: 1px solid black; padding: 2px;">Minimum Documentation Searched⁷</div> <table style="width: 100%; border-collapse: collapse;"> <tr> <th style="width: 20%; border: 1px solid black; padding: 2px;">Classification System</th> <th style="border: 1px solid black; padding: 2px;">Classification Symbols</th> </tr> <tr> <td style="border: 1px solid black; padding: 5px; vertical-align: top;">IPC5</td> <td style="border: 1px solid black; padding: 5px; vertical-align: top;">C 08 G; C 08 J</td> </tr> </table> <div style="text-align: center; border: 1px solid black; padding: 2px;">Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in Fields Searched⁸</div>			Classification System	Classification Symbols	IPC5	C 08 G; C 08 J								
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III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹ <table border="1" style="width: 100%; border-collapse: collapse;"> <thead> <tr> <th style="width: 10%; padding: 2px;">Category¹⁰</th> <th style="width: 60%; padding: 2px;">Citation of Document,¹¹ with Indication, where appropriate, of the relevant passages¹²</th> <th style="width: 30%; padding: 2px;">Relevant to Claim No.¹³</th> </tr> </thead> <tbody> <tr> <td style="text-align: center; vertical-align: top; padding: 5px;">A</td> <td style="padding: 5px;">EP, A1, 0354159 (THE GOODYEAR TIRE & RUBBER COMPANY) 7 February 1990, see the whole document</td> <td style="text-align: center; vertical-align: top; padding: 5px;">1-6</td> </tr> <tr> <td style="text-align: center; vertical-align: top; padding: 5px;">A</td> <td style="padding: 5px;"> <div style="text-align: center;">--</div> EP, A2, 0335819 (THE GOODYEAR TIRE & RUBBER COMPANY) 4 October 1989, see column 13, line 34 - line 57 <div style="text-align: center;">--</div> </td> <td style="text-align: center; vertical-align: top; padding: 5px;">1-6</td> </tr> <tr> <td style="text-align: center; vertical-align: top; padding: 5px;">A</td> <td style="padding: 5px;"> <div style="text-align: center;">--</div> EP, A2, 0284544 (THE GOODYEAR TIRE & RUBBER COMPANY) 28 September 1988, see the whole document <div style="text-align: center;">--</div> </td> <td style="text-align: center; vertical-align: top; padding: 5px;">1-6</td> </tr> </tbody> </table>			Category ¹⁰	Citation of Document, ¹¹ with Indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³	A	EP, A1, 0354159 (THE GOODYEAR TIRE & RUBBER COMPANY) 7 February 1990, see the whole document	1-6	A	<div style="text-align: center;">--</div> EP, A2, 0335819 (THE GOODYEAR TIRE & RUBBER COMPANY) 4 October 1989, see column 13, line 34 - line 57 <div style="text-align: center;">--</div>	1-6	A	<div style="text-align: center;">--</div> EP, A2, 0284544 (THE GOODYEAR TIRE & RUBBER COMPANY) 28 September 1988, see the whole document <div style="text-align: center;">--</div>	1-6
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IV. CERTIFICATION <table style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 50%; border: 1px solid black; padding: 5px;"> Date of the Actual Completion of the International Search 4th November 1991 </td> <td style="width: 50%; border: 1px solid black; padding: 5px;"> Date of Mailing of this International Search Report 26. 11. 91 </td> </tr> <tr> <td style="border: 1px solid black; padding: 5px;"> International Searching Authority EUROPEAN PATENT OFFICE </td> <td style="border: 1px solid black; padding: 5px;"> Signature of Authorized Officer <div style="display: flex; align-items: center;"> <div style="border: 1px solid black; padding: 2px 10px; margin-right: 10px;">M. PEIS</div> <div style="font-family: cursive; font-size: 1.2em;">M. Peiz</div> </div> </td> </tr> </table>			Date of the Actual Completion of the International Search 4th November 1991	Date of Mailing of this International Search Report 26. 11. 91	International Searching Authority EUROPEAN PATENT OFFICE	Signature of Authorized Officer <div style="display: flex; align-items: center;"> <div style="border: 1px solid black; padding: 2px 10px; margin-right: 10px;">M. PEIS</div> <div style="font-family: cursive; font-size: 1.2em;">M. Peiz</div> </div>								
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III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category *	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No
A	<p>US, A, 4064112 (ROTHER ET AL) 20 December 1977, see examples 1-18</p> <p style="text-align: center;">-- -----</p>	1-6

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
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EP-A1- 0354159	07/02/90	AU-B-	611353	06/06/91
		AU-D-	3647489	08/02/90
		JP-A-	2075627	15/03/90
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